

North Carolina Department of Health and Human Services

NC Medicaid

Cystic Fibrosis Medications PA Request Form

(Kalydeco, Orkambi, Symdeko, and Trikafta)

Beneficiary Information

1. Beneficiary Last Name: _____ 2. First Name: _____
3. Beneficiary ID #: _____ 4. Beneficiary Date of Birth: _____ 5. Beneficiary Gender: _____

Prescriber Information

6. Prescribing Provider NPI#: _____
7. Requester Contact Information Name: _____ Phone #: _____ Ext: _____

Drug Information

8. Med requested: _____ 9a. Strength: _____ 9b. Quantity per 30 days _____
9c. Requested Duration (circle # days): 30 60 90 120 180

Requests for KALYDECO (answer questions 10- 14)

10. Does the beneficiary have a diagnosis of Cystic Fibrosis? Yes ___ No ___
11. Is the beneficiary age 6 months or greater? Yes ___ No ___
12. Does the beneficiary have a documented mutation in the CFTR gene that is responsive to ivacaftor? Yes ___
No ___ Please list: _____
13. Is the total daily dose 300mg/day or less? Yes ___ No ___
14. Have a baseline ALT and AST been assessed prior to beginning therapy? Yes ___ No ___

Requests for ORKAMBI (answer questions 15- 19)

15. Does the beneficiary have a diagnosis of Cystic Fibrosis? Yes ___ No ___
16. Is the beneficiary age 2 years or greater? Yes ___ No ___
17. Is there documentation that indicates the beneficiary is homozygous for the *F508del* mutation in the *CFTR* gene?
Yes ___ No ___
18. Is the total daily dose lumacaftor 400 mg/ivacaftor 250 mg or less? Yes ___ No ___
19. Have a baseline ALT and AST been assessed prior to beginning therapy? Yes ___ No ___

Requests for SYMDEKO (answer questions 20- 24)

20. Does the beneficiary have a diagnosis of Cystic Fibrosis? Yes ___ No ___
21. Is the beneficiary age 6 years or greater? Yes ___ No ___
22. Is there documentation that indicates the beneficiary is documented as homozygous for the *F508del* mutation in the *CFTR* gene or does the beneficiary have one mutation in the *CFTR* gene that is responsive to tezacaftor/ivacaftor? Yes ___ No ___
23. Is the daily dose one tablet in the morning and one tablet in the evening? Yes ___ No ___
24. Have a baseline ALT and AST been assessed prior to beginning therapy? Yes ___ No ___

Requests for TRIKAFTA (answer questions 25- 30)

25. Does the beneficiary have a diagnosis of Cystic Fibrosis? Yes ___ No ___
26. Is the beneficiary age 12 years or greater? Yes ___ No ___
27. Is there documentation that indicates the beneficiary has at least one copy of the *F508del* mutation in the *CFTR* gene? Yes ___ No ___
28. Is the total daily dose two tablets (elexacaftor 100 mg, tezacaftor 50 mg and ivacaftor 75 mg) in the morning and one tablet (ivacaftor 150 mg) in the evening or less? Yes ___ No ___
29. Have a baseline ALT, AST, and bilirubin been assessed prior to beginning therapy? Yes ___ No ___
30. If the beneficiary is less than 18 years of age, has a baseline ophthalmic examination been performed? Yes ___ No ___

Signature of Prescriber: _____ Date: _____

(Prescriber signature mandatory)

I certify that the information provided is accurate and complete to the best of my knowledge, and I understand that any falsification, omission, or concealment of material fact may subject me to civil or criminal liability.

This form can be uploaded into the secure NCTracks Provider Portal, faxed, or mailed to NCTracks. Fax all forms and lab work to NCTracks at: (855) 710-1969. Pharmacy PA Call Center: (866) 246-8505.